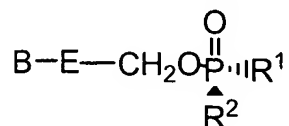


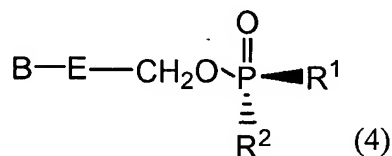
Claim Amendments

Claims 1 – 33 (canceled)

Claim 1 (new) A diastereomerically enriched compound having the structure (3)



which is substantially free of the diastereomer (4)



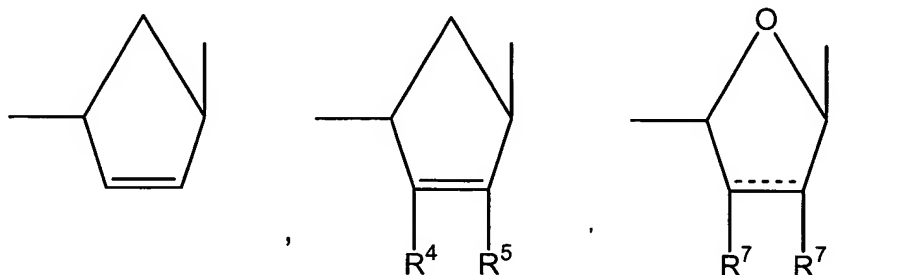
wherein

R^1 is an oxyester which is hydrolyzable *in vivo*, or hydroxyl;

B is a heterocyclic base;

R^2 is hydroxyl, or the residue of an amino acid bonded to the P atom through an amino group of the amino acid and having each carboxy substituent of the amino acid optionally esterified, but not both of R^1 and R^2 are hydroxyl;

E is $-(\text{CH}_2)_2-$, $-\text{CH}(\text{CH}_3)\text{CH}_2-$, $-\text{CH}(\text{CH}_2\text{F})\text{CH}_2-$, $-\text{CH}(\text{CH}_2\text{OH})\text{CH}_2-$, $-\text{CH}(\text{CH}=\text{CH}_2)\text{CH}_2-$, $-\text{CH}(\text{C}\equiv\text{CH})\text{CH}_2-$, $-\text{CH}(\text{CH}_2\text{N}_3)\text{CH}_2-$,



$-\text{CH}(\text{R}^6)\text{OCH}(\text{R}^6)-$, $-\text{CH}(\text{R}^9)\text{CH}_2\text{O}-$ or $-\text{CH}(\text{R}^8)\text{O}-$, wherein the right hand bond is linked to the heterocyclic base;

the broken line represents an optional double bond;

R^4 and R^5 are independently hydrogen, hydroxy, halo, amino or a substituent having 1-5 carbon atoms selected from acyloxy, alkyoxy, alkylthio, alkylamino and dialkylamino;

R^6 and R^6 are independently H, C_1 - C_6 alkyl, C_1 - C_6 hydroxyalkyl, or C_2 - C_7 alkanoyl;

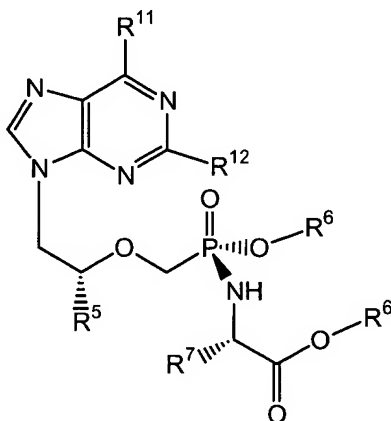
R^7 is independently H, C_1 - C_6 alkyl, or are taken together to form -O- or - CH_2 -;

R^8 is H, C_1 - C_6 alkyl, C_1 - C_6 hydroxyalkyl or C_1 - C_6 haloalkyl; and

R^9 is H, hydroxymethyl or acyloxymethyl;

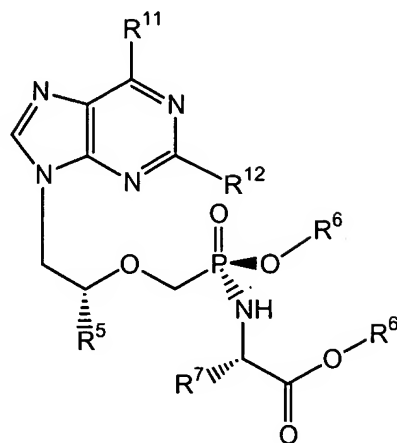
and their salts, free base, and solvates.

Claim 2 (new) A diastereomerically enriched compound having the structure (5a)



(5a)

which is substantially free of diastereomer (5b)



(5b)

wherein

R⁵ is methyl or hydrogen;

R⁶ independently is H, alkyl, alkenyl, alkynyl, aryl or arylalkyl, or R⁶ independently is alkyl, alkenyl, alkynyl, aryl or arylalkyl which is substituted with from 1 to 3 substituents selected from alkylamino, alkylaminoalkyl, dialkylaminoalkyl, dialkylamino, hydroxyl, oxo, halo, amino, alkylthio, alkoxy, alkoxyalkyl, aryloxy, aryloxyalkyl, arylalkoxy, arylalkoxyalkyl, haloalkyl, nitro, nitroalkyl, azido, azidoalkyl, alkylacyl, alkylacylalkyl, carboxyl, or alkylacylamino;

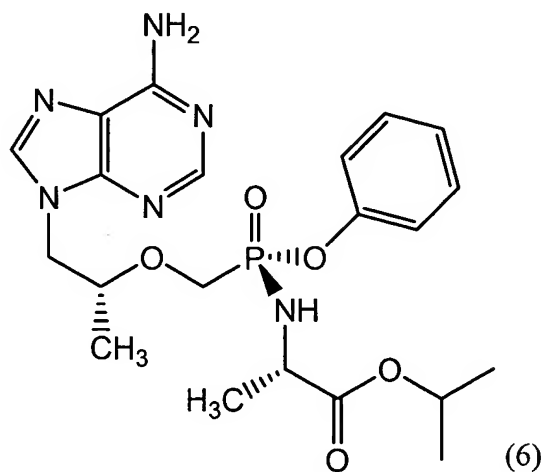
R⁷ is the side chain of any naturally-occurring or pharmaceutically acceptable amino acid and which, if the side chain comprises carboxyl, the carboxyl group is optionally esterified with an alkyl or aryl group;

R¹¹ is amino, alkylamino, oxo, or dialkylamino; and

R¹² is amino or H;

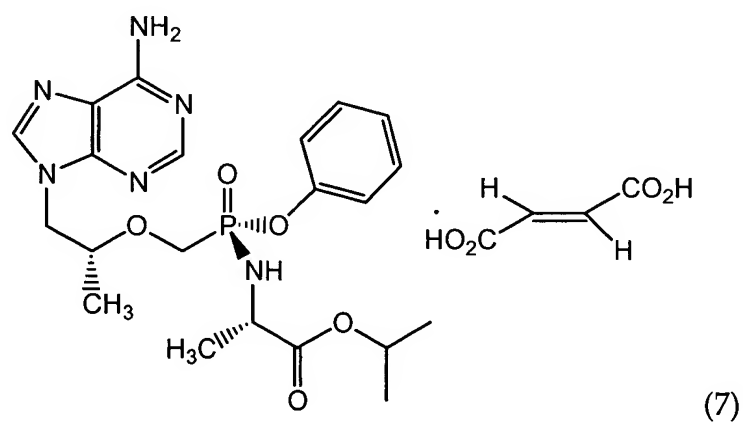
and its salts, tautomers, free base and solvates.

Claim 3 (new) A diastereomerically enriched compound of structure (6)

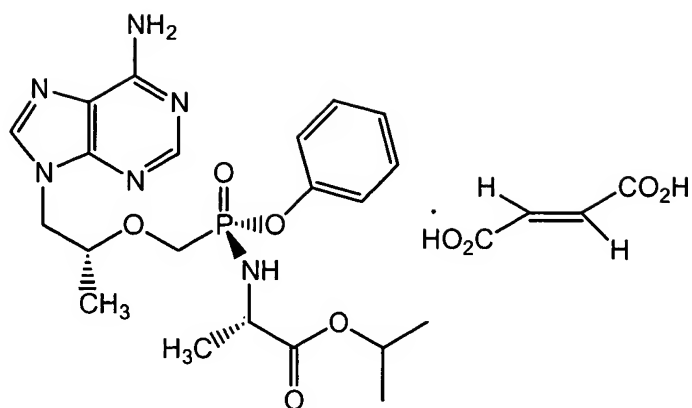


and its salts, tautomers, free base and solvates

Claim 4 (new) A diastereomerically enriched compound of structure (7)



which is substantially free of diastereomer (7a)



(7a)

Claim 5 (new) A composition comprising a compound of any of claims 1-4 and a pharmaceutically effective excipient.

Claim 6 (new) The composition of claim 5 wherein the excipient is a gel.

Claim 7 (new) The composition of claim 5 which is suitable for topical administration.

Claim 8 (new) A method for antiviral therapy or prophylaxis comprising administering a compound of any of claims 1-4 in a therapeutically or prophylactically effective amount to a subject in need of such therapy or prophylaxis.